

**Avian cell lines useful for the production of
substances of interest**

5 The present invention relates to a method for producing
avian cell lines, in particular avian stem cells,
comprising progressive or total withdrawal of growth
factors, serum and/or feeder layer. These spontaneously
established lines are adherent or nonadherent cells
10 capable of proliferating indefinitely in a basic
culture medium. The invention also relates to the cells
derived from such lines which are particularly useful
for the production of vaccines and of substances of
interest.

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Stem cells are cells identified by their culture *in*
vitro from an embryo, from part of an embryo or even
from an adult tissue. The expression stem cell is
understood to mean any pluripotent cell of embryonic or
20 adult origin which has a capacity for self-renewal and
is capable of giving specialized differentiated cells.
In other words, any noncancerous cell capable of
dividing indefinitely in culture and of giving a
daughter cell having the same capacity for
25 proliferation and differentiation as the mother cell
from which it is derived. These isolated cells exhibit
particular morphological and immunocytochemical
characteristics. It is also possible to distinguish the
notion of:

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- embryonic stem cells (CES cells), stem cells
which have the characteristic feature of being
obtained from culturing parts or all of a very
early embryo (blastula stage). These CES cells
35 exhibit *in vitro* all the characteristics of a
stem cell, and *in vivo* the unique capacity of
contributing to the morphogenesis of an embryo
and of participating in germline colonization

when they are reimplanted in any manner whatsoever in a recipient embryo.

- 5 - somatic stem cells (SSC), cells which have all
the characteristics of stem cells when they are
cultured *in vitro*, but which, unlike CES cells,
do not have the potential to colonize *in vivo*
the gonads after infection into an embryo. They
contribute solely to the morphogenesis of the
10 somatic tissues in the embryo.

Unlike already differentiated primary cells, stem cells do not exhibit an easily identifiable characteristic state of morphological differentiation (fibroblasts,
15 adipocytes, macrophage, and the like), but are rather characterized by a state of proliferation and of nondifferentiation. This state results in different behaviors such as a rapid proliferation *in vitro*, a characteristic morphology, the presence of different
20 markers, variable requirements for growth factors and an ability to respond to particular stimuli for induction of differentiation. They are not sensitive to replicative senescence, a critical period for a large number of differentiated primary cells, including the
25 fibroblasts for example.

To maintain avian stem cells *in vitro* for long periods of time, it is necessary to observe specific culture and maintenance conditions as described in Pain et al.,
30 1996; US 6,114,168 and EP 787 180.

The *in vitro* culture of a primary cell under satisfactory medium and growth factor conditions allows it to proliferate only for a certain number of
35 passages. This cell can be obtained directly from a dissociated tissue or from part of this tissue. This number of passages is nevertheless highly dependent on the animal species considered, on the origin of the tissue, on the age of the donor organism and the like.

In most cases, the cell proliferation observed *in vitro* slows progressively and then the cells stop proliferating.

5 This arrest often corresponds to a replicative senescence, known by the term Hayflick limit. This stoppage is thought to be the result of the action of a true molecular clock of which one of the key components is thought to be the length of the telomeres. The
10 telomeres are repeat sequences situated at the end of the chromosomes. The shortening of these repetitive nucleotide structures is the consequence of the replication of DNA on a semiconservative mode. In the absence of the telomerase enzyme, which is in charge of
15 adding the repeat sequences at the end of the chromosomes, a point of no-return is reached with regard to the size of the telomeres, a point beyond which an as yet unknown molecular mechanism for activation of genes involved in controlling the cell
20 cycle is triggered. The cells are then thought to be blocked in the G1 phase in their divisions and are thought to stop proliferating. Numerous factors appear to be involved in this negative control of the cell cycle such as various cyclins, specific kinases, RB and
25 P53 proteins, specific transcription factors such as E2F and many others (mdm2, BTG, p21, and the like). Conversely, the telomerase enzyme can therefore be viewed as a central factor in cell immortality because it maintains the length of the telomeres and therefore
30 makes the cell insensitive to this loss caused by successive divisions.

In an organism under development and during the life of this organism, only a few cell types, including certain
35 lymphocytes, exhibit a permanent expression of telomerase. This activity also appears to be one of the characteristics of stem cells, both at the somatic level (SSC) and at the germline level. This property of expressing, of maintaining of expression and of

"awakening" expression of the telomerase activity is also often associated with the immortal character of a cell maintained *in vitro*. To date, numerous cancer cells are also detected positive for telomerase activity. This activity is thought to be partially responsible for the capacity for uncontrolled proliferation of tumor cells *in vivo*.

This telomerase activity is, in all cases, an excellent marker for the stem cell character and for the germline lineage and for the capacity of a cell to become immortal. Two criteria are therefore used: the telomerase activity and the size of the telomeres.

In an establishment perspective and very briefly, according to the literature and the results already available in many laboratories, the establishment of cell lines may be carried out according to two routes: a spontaneous establishment resulting from noninduced intrinsic genetic damage or a triggered establishment, induced by the use of viruses, retroviruses or by other means such as chemical agents, irradiation, UV (ultraviolet) radiation, and the like. In mammals, for example, the establishment of rodent (mouse, rat, and the like) cells is recognized as being fairly easy spontaneously; on the other hand, the situation is quite different for human cells regardless of their tissue origin (Smith and Pereira-Smith, 1996).

Thus, when it is desired to obtain cell lines from the abovementioned avian stem cells so as to produce a mass of substances of interest *in vitro*, the problem is to be able to maintain cells in culture in an economical medium while avoiding stumbling blocks such as cellular differentiation and senescence.

At the avian level, it is generally accepted that the establishment of primary cells is a rare, or even practically nonexistent event. The only notable

published exception appears to be the DF-1 line which results from the immortalization, described as being spontaneous, of chicken fibroblasts (Foster et al., 1991; patent US 5,672,485, ATCC No. CRL 12203). At the
5 level of this line, recent articles mention the first components of observed deregulation (Kim et al., 2001a; Kim et al., 2001b).

In the immortalization process, a first step leads the
10 proliferating cell to the Hayflick limit which, depending on the cell types, is between 10 and 50 passages. A first spontaneous mutational event then takes place which allows the cell to cross this first blockage, an event which often affects the p53 and pRb
15 genes, and the like. The cells therefore continue to proliferate until the moment when a second blockage occurs, which is in general lifted by new mutations in other genes and by the activation of telomerase, which is often observed.

20 At the avian level, it is in general accepted that the establishment of immortal primary cells is a practically nonexistent event. Accordingly, a large number of lines have been obtained by culturing tumor
25 cells, often directly taken from a biopsy of the tumor. This obtaining *in vitro* is in fact the result of the impairment *in vivo* of certain genes, which are responsible for the appearance of the tumor. An example is provided by the fibroblast line SB-CEV-1, which is
30 isolated from the culturing of a visceral tumor from an animal (ATCC No. CRL 10497, US patent No. 5,846,527). This approach is greatly facilitated by the existence of a very large number of avian viruses and retroviruses which have been identified, isolated and
35 often characterized at the molecular level. Being often oncogenic by their direct or indirect action (activation of a transforming endogenous gene during their integration, expression of the oncogenic protein(s) endogenous to the virus), these viruses

cause tumors and lesions; thus, the obtaining of cell lines in different differentiated lineages is possible from the culturing of the infected organs of the animals. The *in vitro* use of these viruses and retroviruses, isolated *in vivo*, has been developed in addition. There may thus be mentioned nonexhaustively:

- 10 - the lymphoblastoid lines DT40 and DT95, obtained in the presence of the avian leukosis virus (ALV) and in which the myc locus is activated (Baba et al., 1985, ATCC No. CRL 2111, CRL 2112),
- 15 - the turkey lymphoblastoid line MDTC-RP19, established with the Marek's disease virus (US No. 4,388,298, ATCC No. 8135),
- 20 - the lymphoblastoid line ConA-C1, established with the REV virus (reticuloendothelial virus, ATCC No. 12135, US No. 5,691,200),
- 25 - the myelomonocytic line BM2 established with the MH2 virus (Liu et al., 1977, US patent No. 5,388,680),
- 30 - and a whole series of hematopoietic lines obtained with different viruses,
 - o the erythroblastic line HD4 (6C2), obtained with the AEV virus
 - o the monocytic line HD11 (Beug et al., 1979), obtained with the MC29 virus
 - 35 o the granulocytic line HD13 (Golay et al., 1988), obtained with the E26 virus
 - o the mixed hematopoietic line IID57 (Metz and Graf, 1991) also obtained with the E26 virus.

However, the problem in these transformation approaches is the obtaining of cells which produce viruses and carry the activated genome of the viruses and retroviruses used. These activations and this viral presence are brakes on their industrial use as replication support for viruses of interest or for the production of specific proteins under optimum safety conditions.

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Another approach to overexpression of oncogenes, of immortalizing genes (adenovirus E1A gene, polyoma SV40 "large T", and the like) or gene fragments has also made it possible to obtain lines from already differentiated primary cells. These components may be introduced into the cells by simple transfection of a vector allowing the expression of the immortalizing part, but may also be introduced via viruses or retroviruses which have been genetically modified to express these immortalizing components. The origin of the immortalizing components may be avian or otherwise, viral or otherwise. The tropism for avian cells can in fact be linked to the original virus or can also be modified. By way of example, the duck fibroblast line TDF-2A is thus obtained by introducing a first immortalizing gene and then an antiapoptotic gene (Guilhot et al., 1993, patent US 6,255,108). Other methods have been developed, such as the overexpression of p53 (Foster et al., US patent No. 5,830,723).

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In addition, the action of chemical carcinogens directly on the animal according to different modes of administration has allowed, inter alia, the obtaining

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- of the QT6 and QT35 lines, of quail fibroblasts (Moscovici et al., 1977, ATCC No. CRL 1708),
- of the chicken hepatocyte line LMH (Kawaguchi et al., 1989, ATTC No. CRL 2117),

- of the chicken fibroblast line CHCC-OU2 (ATCC No. CRL 12302, US patent No. 5,989,805).

5 The expression immortalization event is understood to mean various actions such as:

- 10 - the action of oxidative, heat or chemical stress capable of inducing modifications in the physiology of the cells and/or mutations,
- 15 - the action of the products of specific genes in the physiology of the cells, such as certain immortalizing genes (oncogenes, protooncogenes, cell cycle genes, antiapoptotic genes and the like),
- 20 - targeted destruction by functional recombination or inactivation of antioncogenes, apoptotic genes, tumor suppressor genes, antiproliferative genes, leading to a functional deregulation of the cell cycle or of the physiology of the cell,
- 25 - the control of proliferation genes by their functional blocking, and the like,
- the action of rays (UV, gamma, X, and the like),
- 30 - the action of chemical mutagens (substances which damage DNA, substances similar to growth factors, and the like),
- 35 - the conjugated action of these various actions taken separately.

In the context of the invention, it has been found that the withdrawal of growth factors, serum and/or feeder

layer leads to the isolation of populations of stem cells, in particular of somatic stem cells, which can grow indefinitely in basic culture media.

5 In addition, apart from the hematopoietic stem cells which are for the most part nonadherent cells, the majority of the cells, obtained according to the prior art techniques mentioned above (fibroblasts, hepatocytes, and the like), exhibit an adherent
10 phenotype. Now, the industrial use of cells, as viral replication support, favors nonadherent cells. This phenotype is advantageous both because of ease of handling which avoids the use of a proteolytic enzyme for dissociation and for the high densities reached by
15 nonadherent cells cultured *in vitro*.

The present invention describes the production of lines which can become spontaneously nonadherent and for which the nonadherence is obtained by a withdrawal of
20 the feeder layer. Because of their growth in suspension, these lines are perfectly suitable for industrial use for the production of substances of interest in bioreactors.

25 In addition to their properties of growing on a basic culture medium, it has been discovered that these cell lines allow the replication of certain viruses in yields equivalent to or even higher than the yields obtained with current methods, which makes these cells
30 particularly useful for the mass production of vaccines.

Description

35 Thus, in a first aspect, the present invention relates to a method for producing avian cell lines, characterized in that it comprises the following steps:

a) culturing avian cells in a medium containing all the factors allowing their growth and an inactivated feeder layer,

5 b) passage by modifying the culture medium so as to obtain progressive or total withdrawal of said factors, of the serum and/or of the feeder layer,

c) establishing adherent or nonadherent cell lines
10 capable of proliferating in a basal medium in the absence of exogenous growth factors, serum and/or inactivated feeder layer.

In the context of the invention, the expression
15 "establishment of a line" is understood to mean maintaining cells in culture *in vitro* over a considerable period of time. Advantageously, the cells derived from the lines obtained in step c) are capable of proliferating for at least 50 days, 100 days, 150
20 days, 300 days or preferably at least 600 days. The 600 days do not constitute a time limit because the cell lines obtained are still alive after much longer time periods. Hence, these lines are considered as being able to grow indefinitely in a basic culture medium
25 free of exogenous growth factors, serum and/or inactivated feeder layer. The expression "line" is understood to mean any population of cells capable of proliferating indefinitely in culture *in vitro* while retaining to a greater or lesser degree the same
30 morphological and phenotypic characteristics.

The cells derived from the lines according to the invention may be avian stem cells, in particular avian somatic stem cells.

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The stem cells according to the invention can serve to obtain differentiated cell lines. Indeed, these stem cells have the property of being pluripotent, that is to say that they have the potential to be induced in

multiple differentiation pathways which can be characterized by various specific markers.

5 These cells can also be precursor cells, which correspond to the partially differentiated cells of an adult or embryonic tissue, by contrast to a stem cell and which is capable of dividing and of giving more differentiated cells. The expression "differentiated cell" is understood to mean any specialized cell of an adult or embryonic tissue, having specific markers or fulfilling specific physiological functions. It is possible, in a particular aspect of the invention, in particular for particular isolates or clones derived from a particular isolate obtained during establishment, for these stem cells to contribute to the germline. In this case, these stem cells established as lines are thought to be embryonic stem cells.

20 Of course, the method mentioned above makes it possible to obtain cellular clones derived from cells obtained from established lines. These clones are cells which are genetically identical to the cell from which they are derived by division.

25 In a particular embodiment, the invention relates to a method as defined above, in which the established lines are adherent stem cells which proliferate in the absence of inactivated feeder layer.

30 In this regard, in the method described above, step b) consists in a withdrawal of the components of the medium (growth factors alone or serum alone or growth factors and then serum or alternatively serum and then growth factors). We also found that it is possible to obtain non adherent cells after several passages, at any moment, from these adherent stem cells that proliferate with or without feeder layer.

In another embodiment, the invention relates to a method as defined above in which the established lines are nonadherent stem cells which proliferate in suspension in a medium free of exogenous growth factors.

In this regard, in the method described above, step b) consists in a progressive or total withdrawal of the feeder layer and then optionally in a withdrawal of the other components of the medium (growth factors and serum).

In another embodiment, the invention relates to a method as described above in which the established lines are nonadherent stem cells which proliferate in suspension in a medium free of serum (serum-free medium).

In another embodiment, the invention relates to a method as defined above, in which the established lines are nonadherent stem cells which proliferate in suspension in a medium free of exogenous growth factors and serum.

In another alternative, step b) consists in a progressive or total withdrawal of the growth factors, optionally followed by a progressive withdrawal of the serum.

In another alternative, step b) consists in a progressive or total withdrawal of the growth factors and/or serum, optionally followed by a withdrawal of the feeder layer.

In addition, the established lines may be cells which proliferate in a serum-depleted medium, in particular in a medium free of serum. The expression serum-depleted is understood to mean a gradual reduction of the concentration of serum spread out over time. This

method allows a selection of clones which adapt to these new, increasingly drastic conditions until stable lines are obtained which are capable of growing in a serum-depleted medium or in a medium completely free of
5 serum.

The method described above may additionally comprise a step in which the cells obtained in step c) are subjected to a selection in culture media used for
10 large-scale production so as to obtain clones suitable for the production of vaccines intended for human or animal therapy.

The cells according to the invention have at least one
15 of the following characteristics:

- a high nucleocytoplasmic ratio,
- an endogenous alkaline phosphatase activity,
- 20 - an endogenous telomerase activity,
- a reactivity with specific antibodies selected from the group of antibodies SSEA-1 (TEC01),
25 SSEA-3, and EMA-1.

Preferably, the cells of the invention have all the abovementioned characteristics.

30 In an additional aspect, the invention relates to a method for producing avian lines, which is mentioned above, in which the cells derived from the lines obtained in step c) are modified in order to allow a better use *in vitro* such as the extension of the
35 greater life span or growth densities or alternatively of the lower nutrient requirements.

Advantageously, the cells derived from established lines are modified in order to produce a substance of

interest, in particular a polypeptide of interest, an antibody or an attenuated virus. Said cells may be modified by any technique accessible to persons skilled in the art, in particular homologous, directed and/or conditional recombination (Cre-Lox or FLP-FRT system), by transformation with any vector, plasmid, in particular with the aid of retroviruses.

The medium used in step a) may comprise at least one factor selected from cytokines, in particular LIF, IL-11, IL-6, IL-6R, CNTF, Oncostatin and other factors such as SCF, IGF-1 and bFGF.

In addition, the inactivated feeder layer used in step a) is preferably composed of fibroblasts, including mouse fibroblasts established as a line. Among these fibroblasts are in particular the STO cells which may or may not be modified or transfected with expression vectors (Pain et al., 1996). In this method, the cells used in step a) are cells obtained by suspending cells obtained from blastodermal disks of fertilized eggs in a culture medium comprising at least one cytokine, b-FGF, and SCF. Said cells are inoculated into a layer of feeder cells, incubated, and then collected.

Step b) consists in a progressive withdrawal of each growth factor added to the medium in step a), in particular a cytokine, b-FGF, and SCF, comprising a passage in a new medium free of at least one of said factors and in repeating various successive passages until the medium is free of all of said factors. The expression progressive withdrawal is understood to mean a removal factor by factor from the culture medium. Alternatively, it is possible to carry out a drastic or total withdrawal, that is to say the removal of all of said factors all at once. Thus, the withdrawal of step b) may consist in progressively reducing the concentration of one or more factors or in culturing the avian stem cells directly in a medium free of one

or more factors or alternatively free of all of said factors.

Step b) may also comprise the withdrawal of the serum.

5 In this regard, the withdrawal may be progressive, by reducing the serum concentration during each passage, for example on passing from 10% to 7.5% and then 3.75% and 2%, tending toward 0% (serum-free medium). Alternatively, a drastic withdrawal may be carried out.

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Step b) may also comprise the withdrawal of the feeder layer. The withdrawal of the feeder layer may also be gradual, by reducing the number of inactivated feeder cells during each passage. Alternatively, it is possible to carry out a drastic withdrawal.

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Of course, the order of withdrawals can vary. For example, it is possible to start with the withdrawal of the growth factors and continue with the withdrawal of the feeder layer.

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Thus, in another aspect, the invention relates to the established cell lines and to the cells derived from said lines which can be obtained from the method described above, said cells being capable of proliferating for at least 50 days, 100 days, 150 days, 300 days, or preferably at least 600 days in a medium free of exogenous growth factor, serum and/or feeder layer.

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These cell lines and the cells derived therefrom are capable of proliferating for at least 50 days, 100 days, 150 days, 300 days, or preferably at least 600 days in a basal medium, in particular in a medium such as DMEM, GMEM, HamF12 or McCoy supplemented with various additives commonly used by persons skilled in the art. Among the additives, there may be mentioned nonessential amino acids, vitamins and sodium pyruvate.

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The invention also relates to the cell lines and the cells derived from such lines described above, characterized in that they are avian stem cells, in particular avian somatic stem cells or avian embryonic stem cells.

These stem cells may be adherent, while proliferating in the absence of the inactivated feeder layer. Alternatively, these stem cells are nonadherent and proliferate in suspension in a basal medium mentioned above.

These cells are also characterized in that they have at least one of the following characteristics:

- a high nucleocytoplasmic ratio,
- an endogenous alkaline phosphatase activity,
- an endogenous telomerase activity,
- a reactivity with specific antibodies selected from the group of antibodies SSEA-1 (TEC01), SSEA-3, and EMA-1.

Advantageously, these cells are genetically modified so as to produce a substance of interest, in particular a polypeptide of interest, an antibody or an attenuated virus.

Cells of the invention can for example support the replication of live or attenuated viruses, in particular the viruses selected from the group of adenoviruses, hepadnaviruses, herpesviruses, orthomyxoviruses, papovaviruses, paramyxoviruses, picornaviruses, poxviruses, reoviruses and retroviruses.

Preferably, the viruses belong to the family of orthomyxoviruses, in particular the influenza virus, to the family of paramyxoviruses, in particular the measles, mumps and rubella viruses.

5 In another embodiment, the viruses replicated on these cells belong to the to the family of poxvirus, in particular canarypox virus, fowlpox virus as well as vaccinia virus.

10 Thus, the invention relates to the cell lines described above, the cells derived from said lines and also the cell lines obtained from cells which have been genetically modified. Preferably, the invention relates to the cell lines derived from step c) of the method
15 described above, characterized in that they are avian stem cells capable of growing indefinitely in a basal medium free of exogenous growth factors, depleted of serum or free of serum and/or of feeder layer.

20 In another aspect of the invention, the cells obtained at the end of step c) may be genetically modified.

The invention also relates to a cell culture comprising cells derived from the cell lines described above, in
25 particular avian stem cells or avian embryonic stem cells, and a basal medium free of exogenous growth factors depleted of serum or free of serum and/or of inactivated feeder layer.

30 In an additional aspect, the invention relates to the use of the cell lines and cells described above for the production of substances of interest, in particular of proteins of therapeutic interest, for the replication of live or attenuated viruses, in particular viruses
35 chosen from the group of adenoviruses, hepadnaviruses, herpesviruses, orthomyxoviruses, papovaviruses, paramyxoviruses, picornaviruses, poxviruses, reoviruses and retroviruses.

Preferably, the cell lines and the cells described above are used for the production of viruses belonging to the family of orthomyxoviruses, in particular the influenza virus, and for the production of viruses
5 belonging to the family of paramyxoviruses, in particular the measles, mumps and rubella viruses.

It is possible to introduce into these lines and cells, used for supporting the replication of live or
10 attenuated viruses, the component(s) necessary for accomplishing the complete viral cycle of the virus so as to obtain, for example, the overexpression of the receptor for the virus at the surface of the cell.

15 Therefore, one best mode of the invention is to use the cells as defined above to produce live or attenuated vaccine, for example recombinant vaccine, comprises culturing the adherent or non adherent cell lines established in step c) according to the process
20 described above, inoculating said cells with viral particles and culturing said cells in a basal medium as mentioned above until cell lysis occurs and newly produced viral particles are released in said medium. The invention has shown to be particularly useful for
25 the production of attenuated virus belonging to the family of poxvirus, in particular canarypoxvirus, fowlpoxvirus and vaccinia virus such as such as native or recombinant vaccinia virus (for example Modified Vaccinia virus Ankara, MVA (such as MVA available under
30 ATCC Number VR-1508) or other orthopoxviruses) and is further described in the examples below.

Furthermore, the invention is aimed at the use of the cells according to the invention for producing
35 recombinant viruses expressing antigens as vaccine against infection diseases such as smallpox and cancer (for example melanoma, prostate cancer, breast cancer, lung cancer, ovary cancer, liver cancer...).

For the remainder of the description, reference will be made to the legend to the figures below.

Legend

5 **Figures 1-3: Growth curves for the cell lines of the invention** (with withdrawal of serum (fig. 2) and with withdrawal of feeder layer (fig. 3)).

10 **Figures 4: Photograph showing the characteristic morphology of avian stem cells**

N: nucleus, n: nucleolus and C: cytoplasm

15 (isolate S86N99, X40 magnification, photograph taken with a Sony Cyber-shot digital camera)

20 **Figures 5: Photograph showing the alkaline phosphatase activity of avian stem cell lines which are adherent or which are in suspension**

25 After fixing (0.1% formaldehyde/0.5% glutaraldehyde, 30 minutes at 4°C), the cells are rinsed twice in 1X PBS and incubated for between 10 and 30 minutes at 37°C in an NBT/BCIP (Nitro Blue Tetrazolium chloride 0.375 mg/ml, 5-bromo-4-chloro-3-indolyl phosphate 0.188 mg/ml, 0.1M Tris pH 9.5, 0.05M MgCl², 0.1M NaCl) solution. The reaction is stopped by two 1X PBS washes and the photographs are taken.

30 **A-** illustrates the characteristic violet coloration of the endogenous alkaline phosphatase activity obtained with the adherent line S86N45 p87, a line cultured with no feeder or factor (X40 magnification, Sony Cyber-shot digital camera).

35 **B-** illustrates the violet coloration characteristic of the endogenous alkaline phosphatase activity obtained with the EB14 line maintained from 8 passages in

suspension, line derived from the S86N45 cells, cultured in suspension with no feeder or factor (X20 magnification, Sony Cyber-shot digital camera).

Example 1: Variable origin of the live material used

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The establishment of cell lines is often greatly linked to the genetic nature of the cell material. Thus, in the murine example, proportionately few genetic bases are permissive to the production of embryonic stem ES
10 cells and often involves a notion of inbred animals. In the case of avian animals, it is difficult to obtain inbred animals for historical reasons and because of the origin of selection of commercial strains; it being of interest precisely to avoid inbreeding. Eggs are the
15 initial source of the cells cultured in this invention. The eggs are preferably used nonincubated, but a few hours of incubation may be necessary in order to obtain the first stages of development of the embryo. The cells obtained are derived from different chicken
20 strains. Among the strains used, there may be mentioned the S86N strain, a commercial strain intended for the production of chicken bearing a quality label, CNRs, the strain intended for the production of chicken bearing a quality label, Marens, a local strain which
25 is genetically and phenotypically well characterized, White Leghorns, a strain more intended for the production of eggs for consumption and a reference strain for research laboratories, and the like. In the latter strain, various origins have been tested
30 including certain eggs (called Valo) obtained from the White Leghorn strain from Lohmann (Germany) considered to be "SPF" (Specific Pathogen Free) eggs kept under very particular health safety conditions. Numerous cell isolates were obtained from various strains, suggesting
35 the general character of the method.

Example 2: Production and establishment of the adherent cells

The eggs are opened, the yolk is separated from the egg white during the opening. The embryos are removed from the yolk either directly or with the aid of a Pasteur
5 pipette, or with the aid of a small absorbent filter paper (Whatmann 3M paper), cut out beforehand in the form of a perforated ring with the aid of a punch. The diameter of the perforation is about 5 mm. These small rings are sterilized using dry heat for about 30
10 minutes in an oven. This small paper ring is deposited on the surface of the yolk and centered on the embryo which is thus surrounded by the paper ring. The latter is then cut out with the aid of small pairs of scissors and the whole removed is placed in a Petri dish, filled
15 with PBS or with a physiological saline. The embryo thus carried away by the ring is cleaned of the excess yolk in the medium and the embryonic disk, thus freed of the excess vitellin, is collected with a Pasteur
pipette.

20 In both cases, the embryos are placed in a tube containing physiological medium (1X PBS, Tris Glucose, medium, and the like). The embryos are then mechanically dissociated and inoculated on a "feeder"
25 into defined culture medium. Among the preferred conditions used for the culturing, preference is given to the culture medium composed of MacCoy medium as basal medium supplemented with fetal calf serum at an initial concentration of 12 to 8%, with nonessential
30 amino acids at 1%, with a mixture of vitamins of commercial origin at 1%, with sodium pyruvate at a final concentration of 1 mM, with beta-mercaptoethanol at a final concentration of 0.2 mM, glutamine at a final concentration of 2.9 mM, with an initial mixture
35 of antibiotics containing gentamycin at a final concentration of 10 ng/ml, penicillin at a final concentration of 100 U/ml and streptomycin at a final concentration of 100 µg/ml. Rapidly after the first passages of the cells, the mixture of antibiotics is no

longer added to the medium. The expression rapidly is understood to mean after the first 3 to 5 passages in general. A mixture of nucleosides may also be added, this mixture being prepared as described above (Pain et al., 1996). Among the basal media tested under these same conditions and which give similar results are the HamF12, Glasgow MEM and DMEM media, the latter supplemented with biotin at a final concentration of 8 mg/l. By way of comparison, the biotin concentration is 0.2 mg/l in the MacCoy medium, 0.0073 mg/l in the HamF12 and 0 in the commercial DMEM and GMEM media.

The growth factors and the cytokines added to the culture medium are preferably factors and cytokines which are recombinant, including mouse SCF at a final concentration of 1 ng/ml, IGF-1 at a final concentration of 1 to 5 ng/ml, CNTF at a final concentration of 1 ng/ml, IL-6 at a final concentration of 1 ng/ml, and the soluble IL-6 receptor at a final concentration of 0.5 ng/ml to 1 ng/ml. In some experiments, some other factors may be added during the first passages. For example up to passage 3 or 10, it is possible to add bFGF to the medium at a final concentration of 1 ng/ml and IL-11 at a final concentration of 1 ng/ml.

The inoculation is carried out into this medium on the inactivated "feeder" composed of mouse fibroblasts established as lines, the STO cells. In some cases, these cells were transfected with simple expression vectors allowing the expression of growth factors such as avian SCF, constitutively in the STO cells. Thus, this "feeder" produces the factor in a form which is soluble and/or attached in the plasma membrane of the cells.

After initial inoculation of the cells directly into this medium, the medium is partially changed the next day, and then partially or completely during subsequent

days, depending on the rate of adhesion observed for the primary cells. After about 4 to 7 days depending on the cases, the initial culture is dissociated and transferred into new dishes in the same initial medium on the inactivated feeder. After three to five passages, the cells are cultured on an inactivated feeder of STO cells which are nontransfected or transfected with an expression vector encoding a resistance to an antibiotic such as the gene for resistance to neomycin, to hygromycin, to puromycin and the like. After about twenty passages, the cells are progressively deprived of growth factors and cytokines. The expression gradual withdrawal is understood to mean a removal factor by factor from the culture medium. Thus, at one passage, SCF is first of all removed, and then, two or three passages later, IGF-1. If the cells do not exhibit morphological alterations or a variation in their average rate of proliferation, the other factors, such as CNTF and IL-6, are then removed. This withdrawal may also be drastic. All the factors are in this case removed all at once. The cells are then observed and are only passaged several days later if their rate of proliferation is modified. The latter solution is generally that which is practiced.

Various isolates are thus obtained and maintained for very long periods of time. The expression very long periods of time is understood to mean periods of the order of several weeks with a minimum of 50 days, preferably periods greater than 200 to 400 days, without limitation in time. Periods greater than 600 days are observed.

Regardless of the support used, all the cells which are adherent are dissociated with a proteolytic dissociation enzyme, such as pronase, collagenase, dispase, trypsin, and the like. Preferably, a proteolytic enzyme of bacterial origin is used in order to avoid any potential contaminant of animal origin.

These cells have the characteristics of embryonic stem cells with a specific morphology illustrated, by way of example, by the **photograph of figure 4** i.e. a small size, a large nucleocytoplasmic ratio, a nucleus with at least one nucleolus which is clearly visible and a very small cytoplasm. These cells are characterized by growth in the form of more or less compact solid masses. The adherent and nonadherent cells exhibit cross-reactivity with a number of antibodies, as described above in Pain et al., 1996 and in patents US 6,114,168 and EP 787 180. The endogenous telomerase activity component is also present and is an important factor in the "stem" nature of these cells.

Cells of different isolates are obtained and maintained for long periods of time.

Table 1 illustrates a few of the characteristics of these isolates

Name	Species	Start	"Stoppage"	Days	Passage	Generation
S86N16	Chicken S86N	26-01-2000	05-08-2001	559	207	692
WL3	Chicken WL	28-06-2000	09-08-2001	403	153	333
Valo4	Chicken Valo	26-09-2000	07-02-2002	401	135	317
S86N45	Chicken S86N	29-01-2001	12-11-2001	287	118	329

It will be noted that the term "stoppage" does not correspond to the end of the proliferation of the cells but to a deliberate stoppage of the cell cultures by the experimenter. The number of generation n is obtained by the formula $X = 2^n$ or X is the theoretical cumulative number of cells. This number is available since the cells are counted at each passage and during each inoculation. The complete history of the culture is thus available.

Example 3: Passage of the cells

One of the characteristics of stem cells, in particular somatic stem cells and embryonic stem cells, is their capacity to proliferate *in vitro* for considerable periods of time. In order to propagate and to passage the cells, the culture medium is changed and replaced with fresh medium a few hours before their passage. The curve presented in **figure 1** illustrates a profile of cell growth and establishment.

Example 4: Doubling time and average division time

Starting with the established cells in culture and the cells presented in the preceding examples, a mean division time can be calculated. For all the independent isolates obtained, the rate of proliferation increases slightly during successive passages, thus causing the average division time during the establishment of the cells to vary. In the adherent phase, the cells are initially inoculated on an inactivated feeder layer and are passaged regularly at a constant initial inoculation density of $1 \text{ to } 2 \times 10^6$ cells per 100 mm dish. Table 2 illustrates the doubling time (d) and the mean division time (MDT in hour) for 3 established cell types as a function of the culture time. It is observed that the mean doubling time decreases during the establishment.

Table 2:

Cells/days	50	100	150	200	250	300	350	400	450	500	550
S86N16 (d)	0.30	0.63	1.00	0.86	1.13	1.15	1.47	1.70	1.94	1.50	1.9
S86N16 (MDT)	80	38	24	27.9	21.2	20.9	16.3	14.1	12.4	16	12.6
S86N45 (d)	0.49	0.89	0.89	1.45	2.15	x	x	x	x	x	x
S86N45 (MDT)	49	26.8	27	16.5	11.1	x	x	x	x	x	x
Valo4 (d)	0.03	0.61	1.00	1.17	1.26	1.03*	1.08*	1.25*	x	x	x
Valo4 (MDT)	>48	39.3	24	20.5	19	23.3	22.2	19.2	x	x	x

The mean doubling time d is established for the period
 5 of time indicated in days with the following formula:

$$d = (1/\log_2 \times (\log X_2/X_1)) \times 1/(T_2-T_1)$$
 where X₂ and X₁
 are total numbers of cells at the times T₂ and T₁. This
 formula is the direct consequence of the calculation of
 the number of generations N by the formula $X = 2^n$
 10 presented in example 1. The mean division time (MDT) is
 then obtained in hours by dividing 24 hours by d.

* The Valo cells are passaged during this establishment
 on a plastic support without the presence of a feeder.
 15 The doubling time decreases and then increases again,
 when the cells become rehabilitated to this new
 environment.

Example 5: Control of the level of serum for the
 20 **proliferation of the lines**

During the obtaining of these lines, the culture media
 used are conventional culture media comprising a base
 (DMEM, GMEM, HamF12, McCoy, and the like) supplemented
 25 with various additives such as nonessential amino
 acids, vitamins, and sodium pyruvate. This complex
 medium comprises fetal calf serum, which remains a
 central component of the culture, even though
 components of different origins, including plant
 30 components, can be gradually used. A process for
 controlling and habituating the cells to relatively low

proportions of fetal calf serum is presented. It is thus possible to maintain cells in high proliferation (division time >1) with low percentages of serum (2% for example in the case of the S86N16 cells).

5

The curves presented in **figure 2** illustrates the relative reduction of serum for a given cell type: S86N16 cells, The doubling time and the mean division times were also calculated and presented in **table 3**. It will be noted that the mean division time increases as a function of the relative reduction in serum. A recovery phase is nevertheless observed after some time in culture under the conditions mentioned. This time remains nevertheless less than 24 h (d>1), which already represents a very advantageous proliferation in industrial terms even at serum concentrations of 2%, which is already relatively low. Improvements with regard to the different metabolites to be used may be envisaged in order to increase this time and still further optimize the culture conditions.

Table 3:

Condition	10%	7.5%	3.75%	2%
d	2.02	1.51	1.47	1.08
MDT	11.9	15.8	16.3	22.2

25 The examples are taken between passages p204 and p179 for the 10% condition, between p198 and p176 for the 7.5%, between p224 and p201 for the 3.75% and between p216 and p199 for the 2%.

30 **Example 6: Deprivation of the cells of feeder layer**

Under the initial culture conditions, the presence of a layer of inactivated cells appears to be necessary in order to obtain embryonic stem cells as was described above. This feeder layer no longer appears to be necessary after a number of passages. Only the "culture

35

treated" plastic appears to be important. Indeed, one of the characteristics of some eukaryotic cells is to proliferate in adherent form. In order to facilitate the adhesion of the cells, the various plastic materials used are "culture" treated. They undergo during their manufacture a treatment which adds charges at the surface of the plastic, which charges promote the adhesions of the extracellular matrix of the cells. By contrast, the cell culture untreated plastic, often called plastic of bacteriological quality, is not surface treated by addition of specific feeders. The adhesion of the cells thereto is generally very difficult, or even impossible, or then induces changes in morphology, and in behavior which are often drastic. This distinction between the two plastic qualities makes it possible to obtain, depending on the inoculations which are carried out therein, cells with different behaviors. Gradual deprivation of the cultures of inactivated "feeder" makes it possible to obtain, after a few passages, homogeneous cultures of stem cells directly inoculated on "culture treated" plastic.

The comparative growth curves for the cells maintained in the presence and in the absence of inactivated "feeder" are presented with the case of the S86N16 cells in **figure 3**. This adaptation of the cells is progressive so as not to lose the stem cell character of the cells initially maintained on a "feeder". Progressive derivatives are thus made. The obtaining of cells which proliferate on plastic is the accomplishment of the withdrawal process. In **table 4**, the division times show sensitivity of the cells to their environment. As in the case of the progressive withdrawal of serum, an adaptation is obtained with a recovering effect on the cells after a few passages under the conditions defined.

Table 4:

Condition	1.2	0.5	0.3	plastic
d	1.95	1.84	1.39	1.42
MDT	12.3	13	17.3	16.9

The examples are taken between the passages p154 and p131 for the 3 conditions 1.2×10^6 , 0.5×10^6 and 0.3×10^6 feeder cells and between p161 and p139 for the condition on plastic alone.

Example 7: Deprivation of the cells in growth factors

Under the initial culture conditions, the presence of growth factors is necessary. It is possible to schematically distinguish two families of factors: the cytokines and the trophic factors.

The cytokines are mainly cytokines whose action is through a receptor which is associated with the gp130 protein. Thus, LIF, interleukin 11, interleukin 6, CNTF, oncostatin and cardiotrophin have a similar mode of action with the recruitment at the level of the receptor of a specific chain and the combination of the latter with the gp130 protein in monomeric or sometimes heterodimeric form. In a few cases, the combination of a soluble form of the receptors, a form described inter alia for the receptors for interleukin 6 and CNTF, makes it possible to increase the proliferative effect observed. It has been previously shown that the addition of at least one of these cytokines appeared to be necessary for obtaining embryonic stem cells.

The trophic factors are mainly SCF, IGF-1 and bFGF, which are also used at the start of the culture, as described above. Their presence is also necessary for obtaining and amplifying the cells.

By progressively reducing these growth factors, it is possible to obtain, after a few passages, culture conditions which allow the proliferation of the embryonic or somatic stem cells without the addition of an exogenous growth factor. The different markers used to characterize these cells are always positive for the cells maintained with no factors.

Example 8: Comparison of the media used

10

Inoculated into different media, the cells are not obtained with the same frequencies. Comparison of the compositions of the media makes the identification of one of the components in particular difficult. It appears more likely that the whole combination allows an improvement in the physiology of the cells. Among the preferred media, the Ham F12 medium, the MacCoy medium, the DMEM medium and a DMEM medium enriched with biotin will be noted. Starting with such an isolate, adaptation trials are carried out in these different media.

20

Example 9: Establishment of the nonadherent cells

During the successive passages of the stem cells, a high-density inoculation directly into the bacteriological dish makes it possible to obtain, after a few passages, embryonic cells which become detached from their substrate and which proliferate in suspension in the form of small regular aggregates. This proliferation is encouraged over several passages by mere dilution, mechanical dissociation and nonuse of proteolytic enzyme. The stirring of the cultures is generally carried out but does not represent a distinguishing factor for obtaining nonadherent cells. Like the adherent cells, these cells have a characteristic morphology of stem cells, i.e. a small size, a large nucleocytoplasmic ratio, a nucleus with at least one nucleolus which is clearly visible and a

30

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very small cytoplasm. These cells are characterized by a growth in small aggregates which are more or less compact. These nonadherent cells exhibit cross-reactivity with a number of antibodies, as described above in Pain et al., 1996. These cells are also positive for the endogenous telomerase activity (as presented in example 10 for the EB1, EB4 and EB5 cells). In a nonadherent phase, the cells exhibit a high proliferation in different media. The initial inoculation density and the very regular supply of fresh medium provides high densities which may range above 1×10^6 cells per ml. **Table 5** summarizes the main characteristics of a few isolates (parental cells, initial passage of the making into a suspension, number of days maintained in culture in suspension, number of passages and of generations obtained before voluntary stoppage of the maintenances). It can thus be noted that the passage for the making into a suspension can vary from one isolate to another (see isolate EB1 and EB14) and the proliferation rate (see isolate EB3 and EB14).

Table 5:

Name	Parental cells	Initial passage	Start	Days	Passages	Generations
EB1	S86N16	p111	20-01-2001	184	41	120
EB3	S86N16	p118	23-01-2001	381	17	40
EB4	S86N45	p100	25-09-2001	44	17	40
EB5	S86N45	p100	25-09-2001	44	17	40
EB14	S86N45	p81	05-09-2002	70	24	65

It will be noted that the term "start" corresponds to the cells being placed under nonadherence.

Example 10: Characterization of the established cells

The stem cells maintained for long culture times are characterized with the same criteria as those described above (Pain et al., 1996). It is thus possible to

regularly detect the endogenous alkaline phosphatase activity, illustrated by the **photograph of figure 5**, the endogenous telomerase activity and reactivity with specific antibodies such as the antibodies SSEA-1
5 (TEC-01) and EMA-1.

One of the important criteria during the establishment of the cells is the presence of telomerase. Various tests were carried out during the maintenance of the
10 cells in culture using a TRAP detection kit (Telomerase PCR Elisa, Roche). The cells are detected positive after various passages in culture. Thus, the telomerase activity is detectable for the S86N16 cells, the S86N45 cells and for the EB1, EB4 and EB5 cells which are
15 derived therefrom in a nonadherent form (see **table 6**). The CEFs (Chicken Embryonic Fibroblasts) maintained in primary culture are considered as negative. The threshold of an OD < 0.2 is the threshold recommended by the kit as the negative threshold. All the analyses
20 were carried out on an equivalent of 2000 cells.

Table 6: Assay of the telomerase activity in various lines at various passages

Cells	Passage	Telomerase OD
S86N16	p12	1.7
	p29	2.8
	p185	0.97
	p204	0.95
S86N16 EB1	p134	1.1
S86N45	p50	0.87
	p58	1.1
	p66	0.96
	p94	1.2
S86N45 EB4	p112	1.4
S86N45 EB5	p112	0.94
CEF*	p4	0.07

Example 11: Transfection and induction of the cells

The stem cells maintained in a growth over the long term are transfected with various expression plasmids.

5 It has been shown that avian stem cells could be transfected (Pain et al., 1996). In particular, the nonadherent cells are transfected and various sorting systems make it possible to identify the stably transfected cells (cell sorting, limiting dilution, and

10 the like). These genetic modifications can be made at the undifferentiated stage of the stem cell. Once this modification has been obtained, the cell is then induced to differentiate spontaneously or by addition of a differentiation inducer. In this case, it is

15 possible to use retinoic acid at concentrations of 10^{-8} M to 10^{-6} M, or dimethyl sulfoxide at concentrations of 1 to 2% final or sodium butyrate at concentrations of 10^{-4} to 10^{-8} M, or phorbol ester (TPA, PMA, and the like) or lipopolysaccharides (LPS) at concentrations of

20 1 to 5 $\mu\text{g/ml}$ final. In another example, the cells can form embryoid bodies in suspension, which embryoid bodies can be caused to adhere to plastic after dissociation or nondissociation of the cells constituting them. These differentiated cells then

25 proliferate but have a more limited capacity for proliferation over the long term. By targeting the genetic modification on a gene which influences the proliferation of the cells, it is possible to make these differentiated cells capable of proliferating

30 over the long term.

Example 12: Infection of the cells

The adherent and nonadherent cells can be infected with

35 different viruses and retroviruses including avian viruses and retroviruses. These cells can thus serve as a replication support for the production of viral stocks intended for the production of live, attenuated or inactivated human and veterinary vaccines depending

on the cases. Among the viruses of interest, there may be mentioned those of the family of adenoviruses (such as Human Adenovirus C, Fowl Adenovirus A, Ovine Adenovirus D, Turkey Adenovirus B), circoviridae (such as Chicken Anemia Virus, CAV), certain coronaviruses, such as avian infectious bronchitis virus (IBV), flaviviruses (such as Yellow fever virus and hepatitis C virus), hepadnaviruses (such as Hepatitis B virus and Avihepadnaviruses such as Duck hepatitis B virus); herpesviruses (such as Gallid herpesvirus, HSV (Herpes simplex virus) and Human herpesvirus 1, 3 and 5), orthomyxoviruses (such as the influenza virus: Influenzavirus A, Influenzavirus B and Influenzavirus C), papovaviruses (such as polyomavirus and more particularly Simian virus 40), paramyxoviruses (such as measles, mumps and rubella viruses and such as respiroviruses and pneumoviruses such as human respiratory syncytial virus and Metapneumovirus such as Avian pneumovirus), picornaviruses (such as polio virus, hepatitis A virus, and such as Encephalomyocarditis virus and foot-and-mouth disease virus), poxviruses (such as fowlpox virus and avipox viruses including Canarypox viruses, Juncopox viruses, Mynahpox viruses, Pigeonpox viruses, Psittacinepox viruses, Quailpox viruses, Sparrowpox viruses, Starlingpox viruses, Turkeypox viruses), orthopoxvirus such as vaccinia virus, MVA, and reoviruses (such as rotaviruses), retroviruses (such as ALV, avian leukosis virus, Gammaretroviruses such as Murine leukemia virus, Lentiviruses such as Human immunodeficiency virus 1 and 2) and Togaviridae such as Rubivirus, in particular Rubella virus.

Example 13: Protocol for infecting a nonadherent avian cell line (EB1) with a virus

Amplification of the cells:

The EB1 or EB14 cells are inoculated into a medium, preferably McCoy's 5A, HAMF12 or DMEM medium, or any

other medium of interest, containing 5% serum at a concentration of 0.2×10^6 cells/ml for an initial volume of 50 ml in general. They are maintained in culture at 39°C and at 7.5% CO₂, with stirring. Fresh
5 medium is added every day for the 3 to 4 days for which the amplification lasts in order to reach a cell concentration of 1 to 3×10^6 cells/ml for a final culture volume of 100 to 250 ml.

The cells in suspension are collected and centrifuged
10 for 10 min at 1 000 rpm approximately. The pellet is resuspended in 20 to 50 ml of 1X PBS (Phosphate buffer Salt). The cells are then counted, centrifuged and the pelleted cells are taken up in a serum-free medium at a final concentration of 3 to 5×10^6 cells/ml. Several
15 tubes are then prepared under these conditions containing 3 to 5×10^6 cells per tube.

Preparation of the virus and infection:

The viral stock having a known titer is rapidly thawed
20 at 37°C and diluted in serum-free medium at a titer of $10 \times$ to $1\,000 \times$ the concentration necessary for the final infection. The cells are infected with the virus of interest at an m.o.i. (multiplicity of infection) of 0.01 to 0.5 according to the types of virus, which
25 involves adding between 0.1 and 10% volume/volume of viral suspension to the cellular pellet. After incubating for 1 hour at an optimum temperature for the virus, in general from 33 to 37°C, the cells are again centrifuged and the medium removed with care. This step
30 is found to be often necessary in order to limit the effect of the initial virus in the subsequent process. One of the possibilities is to directly dilute the cells without centrifuging them again with serum-containing medium (5% of serum) at a final
35 concentration of 0.2 to 1×10^6 cells/ml and incubated again.

Harvesting of the supernatant and of the cells:

After 2 to 4 days of incubation, depending on the viral kinetics and the potential cytopathic effect of certain viruses, the medium containing the cells or the cellular debris is harvested. Depending on the viruses, only the pellet or the supernatant may be of interest and contain the viral particles. The cells are harvested and centrifuged. The collected supernatant is centrifuged again for 5 to 10 minutes at 2 500 rpm, and stored at -80°C before purification of the particles. An aliquot is collected in order to carry out the titration. The cellular pellet is taken up in 5 ml of serum-free medium, sonicated and centrifuged for 5 to 10 minutes at 2 500 rpm. The supernatant obtained is stored at -80°C up to the purification and the titration of an aliquot.

The viral infection and production efficiencies are compared between the various conditions performed. For the viruses with cytopathic effects, the titrations are in general carried out by the lysis plaque technique.

Example 14: Protocol for infecting an adherent avian cell line (S86N45) with a virus

25

Preparation of the cells:

The cells are inoculated 48 hours before the infection into T150 flasks at a concentration of between 0.03 and 0.06×10^6 cells/cm² in a medium, preferably MacCoy's 5A, HAMF12 or DMEM medium, or any other medium of interest, containing 5% serum. They are maintained at 39°C and 7.5% CO₂.

Infection:

35 The viral stock having a known titer is rapidly thawed at 37°C and diluted in serum-free medium at a titer of $10 \times$ to $1\,000 \times$ the concentration necessary for the final infection. The cells are infected with the virus of interest at an m.o.i. (multiplicity of infection) of

0.01 to 0.5 according to the types of virus, which involves adding between 0.1 and 10% volume/volume of viral suspension to the cellular pellet. The infection is generally carried out in a minimum of medium (from 5
5 to 10 ml for a 75 cm² flask) in a medium containing 0% serum.

After incubating for 1 hour at the optimum temperature for the virus, in general from 33 to 37°C, 20 ml of
10 medium 5% are added to the flasks. In a particular case, the cells can be washed with PBS in order to remove the particles which might be attached to the cells. In the case of a cytopathic virus, the cells are observed daily after the infection in order to monitor
15 the appearance of the cell lysis plaque, which indicates good progress of the infection.

Harvesting of the supernatant and of the cells:

After 2 to 4 days of incubation, depending on the viral
20 kinetics and the potential cytopathic effect of certain viruses, the medium containing the supernatant, the cells and the cellular debris are harvested. Depending on the viruses, only the pellet or the supernatant may be of interest and contain the viral particles. The
25 cells are harvested and centrifuged. The collected supernatant is centrifuged again for 5 to 10 minutes at 2 500 rpm, and stored at -80°C before purification of the particles. An aliquot is collected in order to carry out the titration. The cellular pellet is taken
30 up in 5 ml of serum-free medium, sonicated and centrifuged for 5 to 10 minutes at 2 500 rpm. The supernatant obtained is stored at -80°C up to the purification and the titration of an aliquot.

35 The viral infection and production efficiencies are compared between the various conditions performed. For the viruses with cytopathic effect, the titrations are in general carried out by the lysis plaque technique.

Example 15: Replication of a recombinant avipox on nonadherent avian stem cells of the EB1 line

- 5 The nonadherent stem cells EB1 at passage 138 are amplified and then infected at an m.o.i. of 0.1 with a recombinant avipox producing a protein of interest. After the infection, the cells are maintained in a spinner for the 4 days for which the infection lasts.
- 10 An aliquot is removed from the second day and then the next two days in order to monitor the variation of the viral titer both in the supernatant and in the intracellular content, after lysis of the cellular content. The titration is carried out by the lysis
- 15 plaque technique.
- Table 7** illustrates the results obtained. These results demonstrate the very satisfactory replication of the recombinant avipox on the EB1 stem cells. Thus, the infectious titer progresses throughout the culture and
- 20 the course of the infection, reaching a maximum of 7.2 PFU/cell (PFU: Plating Forming Unit) after 4 days of incubation. This titer is at least equivalent to that obtained for this same recombinant avipox on primary chicken embryo cells.
- 25 This titer can be improved by specific culture conditions and optimization procedures.
- It will also be noted that at least equivalent infectious titers were also obtained on a larger scale in 3 liter bioreactors.

Table 7: Kinetics of titration of the recombinant avipox on nonadherent EB1 stem cells

Sampling (h after infection)	50 hours	74 hours	97 hours
Cellular fraction (Log PFU/ml)	6.40	6.37	5.99
Supernatant (Log PFU/ml)	5.56	5.8	6.29
Total (Log PFU/ml)	5.78	5.94	6.31
PFU/cell	2.2	3.2	7.2

5

Example 16: Replication of Modified Vaccinia virus Ankara (MVA) on adherent and nonadherent avian stem cells of the S86N45 line and EB14 line.

- 10 The MVA virus (titer $2,5 \times 10^7$ TCID₅₀/ml in 0.5 ml vials) was received under frozen conditions.
For safety reasons, the MVA virus and infected cells were kept under controlled conditions (-80°C freezer) and the contaminated plastic material was placed into
15 hypochloride solution for more than 1 hour and then place into a bag for full and complete autoclave inactivation.

Experiments and results

- 20 The S86N45 and EB14 cells were thawed in a HamF12 based complete medium.
The adherent S86N45 cells were amplified quite rapidly, with a good growth rate and a nice morphology..
The cells were infected 1 hour in 2 ml of PBS with the
25 different m.o.i. of interest with no washing with PBS after the infection. The medium was just added to the

complete infectious medium, i.e. the added virus was not removed.

After 3 days of infection, the cell lysis appears to be proportional to the used m.o.i. This cytopathic effect is a good indicator of the virus infection of the cells. So cells and supernatant are harvested and stored at - 80°C before purification of particles and/or titration.

10

The non adherent EB14 cells were amplified. The cells were infected, not washed, and the complete medium directly added on the inoculum after 1 hour of contact with viral particles. After 3 days, a characteristic cell lysis was observed. The non infected cells used as the control were counted and a good growth was demonstrated, showing good culture conditions and therefore confirming an efficient lysis by the virus in the infected culture. Cells and supernatant are harvested and stored at - 80°C before purification of particles and/or titration (see table 8).

20

Table 8 : Results of the titration

Cell Type	M.O. I.	Trial	Virus / pellet of 10E6 cells	Titer/mL based on 14ml total volume	Virus yield / cell	Average virus yield / cell
S86N45	0	1	0	0	0	0
	0	2	0	0	0	
	0,1	1	1-3 x 10^7	≥ 6 x 10^5	5-15	5-15
	0,1	2	1-3 x 10^6	1-3 x 10^5	1-5	
	0,01	1	1-3 x 10^7	1-3 x 10^6	15-30	≥ 30
	0,01	2	4-6 x 10^7	1-3 x 10^6	≥ 30	
EB14	0	1	0	0	0	0
	0,1	1	1-3 x 10^7	1-3 x 10^6	15-30	15-30
	0,1	2	1-3 x 10^7	1-3 x 10^6	15-30	

0,05	1	$\geq 6 \times 10^7$	$4-6 \times 10^6$	≥ 50	≥ 50
0,05	2	$1-3 \times 10^7$	$1-3 \times 10^6$	15-30	
0,01	1	$1-3 \times 10^7$	$1-3 \times 10^6$	15-30	15-30
0,01	2	$1-3 \times 10^7$	$1-3 \times 10^6$	15-30	

Applicants hereby incorporate by reference
PCT/FR03/00735 filed on March 7, 2003, and French
application serial no. 0202945 filed March 8, 2002, and
5 all references cited herein.

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